

Attorney Docket No.: MORN-0010
(108347.00021)

PATENT APPLICATION

HIGHLY ACIDIC METALATED MIXTURE OF INORGANIC ACIDS

Inventors: Maurice Clarence Kemp
4401 Woodthrush Drive
El Dorado Hills, California 95762

Robert Blaine Lalum
6949 Greenback Lane #B
Citrus Heights, California 95621

David E. Lewis
816 Third Avenue
Eau Claire, WI 54703

Assignee: Mionix Corp.
1376 Lead Hill Blvd., Suite 130
Roseville, California 95661

T. Ling Chwang
Jackson Walker L.L.P.
2435 N. Central Expressway, Suite 600
Richardson, Texas 75080
Tel: 972-744-2919
Fax: 972-744-2909

HIGHLY ACIDIC METALATED MIXTURE OF INORGANIC ACIDS

BACKGROUND

5 This invention relates to a highly acidic metalated mixture of inorganic acids ("HAMMLA"), to its preparation, and to its uses.

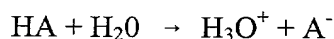
10 In the late 80's and early 90's, researchers in Japan developed strong ionized water ("SIW") as disinfectants. The SIW was established as water with pH 2.7 or less, having an oxidation-reduction potential of 1,000 mv or more, and chlorine concentration of 0.8 ppm or more. The SIW is prepared by electrolysis of water.

Electrolysis of tap water has also been used to produce "strong acid water" and "strong alkali water" both of which were claimed to have antiseptic properties.

15 International Publication WO 94/09798 describes a pharmaceutical composition for treatment of disease, injury and other disorders. The pharmaceutical composition comprises a complex of a calcium-containing component and a sulfate-containing component in a pharmaceutically acceptable carrier. The reference teaches the isolation from natural materials, such as peat, the inorganic compositions. The inorganic preparations comprise
20 an alkaline, aqueous or organic, or mixture thereof, extract of peat. Peat is extracted with aqueous solutions, organic solutions or water-miscible organic solvents at temperature from below room temperature up to the boiling point of the solvents. The preferred extracting solvents are those having a pH of at least 9. Biologically active constituents of fractionated peat preparations were
25 identified as $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$ (gypsum), $\text{CaSO}_4 \cdot \text{K}_2\text{SO}_4 \cdot \text{H}_2\text{O}$ (syngenite, also

referred to as the double salt of gypsum) and $\text{K}_3\text{Na}(\text{SO}_4)_2$ (apthitalite) by X-ray powder diffraction analysis. The reference also describes the synthesis of syngenite.

Chemists describe and measure the ability of a substance to donate protons $[\text{H}^+]$ to a chemical reaction as the pka of that substance where



Although a hydronium ion is usually represented by H^+ or H_3O^+ , its true formula is not certain. The aggregate could be H_5O_2^+ , H_7O_3^+ , or most likely H_9O_4^+ .

The ability to produce large quantities of positively charged water would be extremely important in a large number of applications where water is the medium of the reaction. Positively charged water has the ability to donate protons $[\text{H}^+]$. The donation of a proton is usually an intermediate step in any acid hydrolysis reaction. Acids are the chemical reagent used to donate protons in an aqueous solution.

A strong acid is used to neutralize and remove the lime, or quicklime, from the brick and mortar. A strong acid, such as hydrochloric acid, also known as muriatic acid, is also used to clean hard water spots on shower stalls, windows, glass, toilets, urinals, mirrors and other surfaces. Hydrochloric acid is used to de-scale water towers and heat exchangers and to adjust the pH of the waste water effluent.

A full strength mineral acid, such as hydrochloric acid, is extremely corrosive to many substances, including metals. In addition, hydrochloric acid at a low pH of 0.5 or so will burn a human skin in seconds. The acid is also very harmful in that it emits fumes irritating to mucous membrane. If left near other chemicals, like bleach, hydrochloric acid will interact with them, even through a typical plastic bottle.

The control of microbial growth is necessary in many practical situations, and significant advances in agriculture, medicine and food science have been made through study of this area of microbiology. "Control of growth" means to prevent growth of microorganisms. This control is effected in one of two basic ways: (1) By killing microorganisms; or (2) by inhibiting the growth of microorganisms. Control of growth usually involves the use of physical or chemical agents which either kill or prevent the growth of microorganisms. Agents which kill cells are called "cidal" agents; agents which inhibit the growth of cells, but without killing them, are referred to as "static" agents. Thus the term "bactericidal" refers to killing bacteria and "bacteriostatic" refers to inhibiting the growth of bacterial cells. A "bactericide" kills bacteria, a "fungicide" kills fungi. "Sterilization" is the complete destruction or elimination of all viable organisms in or on an object being sterilized. The object is either sterile or not, there are no degrees of sterilization. Sterilization procedures involve the use of heat, radiation or chemicals, or physical removal of microorganisms.

Microorganisms tend to colonize and replicate on different surfaces resulting in adherent heterogenous microbial accumulations termed "biofilms." Biofilms may form on surfaces of food substances, feed substances, and instrumentations. The microorganisms in the biofilms may include bacteria, fungi, viruses, and protozoans. Since food safety is a national priority, any product that can help by solving a multitude of problems associated with food production is desirable. Removal and control of biofilms which harbor

dangerous microbial contamination is a sanitation goal that needs to be achieved. It is also desirable to be able to safely decontaminate water and nutriment by lowering pH to levels where contaminants would react and organisms cannot live.

5 As used herein, the term “nutriment” means something that nourishes, heals, or promotes growth and repairs the natural wastage of organic life. Thus, food for a human and feed for an animal are all examples of nutriment materials. Other examples of nutriment materials include beverages, food additive, feed additive, beverage additive, food supplement, feed supplement, 10 beverage supplement, seasoning, spices, flavoring agent, stuffing, food dressing, pharmaceutical, biological product, and others. The nutriment materials can be of plant origin, animal origin, or synthetic. Current sanitizing, disinfectants and pesticides products on the market for these uses contain residues of chlorine, ammonia, organic iodine, metal salts and other deleterious residues. It is 15 desirable to have a way that would preclude these residues by promoting acid hydrolysis without the presence of deleterious chemicals. Additionally, this method should generate few hazardous volatile gases. Importantly, it is highly desirable to have a composition that can control and the growth of, and kill, microorganisms and, at the same time, destroy the products generated by, or 20 associated with, the microorganisms.

SUMMARY

One embodiment of the present invention involves a highly acidic metalated mixture of inorganic acids ("HAMMIA"). The composition has an acidic pH, and can be isolated from a mixture prepared by mixing a salt of phosphoric acid, and a preformed, or in-situ generated, solution or suspension of an acidic sparingly-soluble Group IIA complex ("AGIIS"), wherein the solution or suspension of AGIIS is in an amount sufficient to render the acidic pH of the composition to be less than about 2. Another embodiment of the present invention involves a composition having an acidic pH, the composition is isolated from a mixture prepared by mixing a salt of phosphoric acid, and a preformed, or in-situ generated, solution or suspension of AGIIS, wherein the solution or suspension of AGIIS is in an amount in excess of the amount required to completely convert the salt of phosphoric acid to phosphoric acid. Still another embodiment of the present invention involves an adduct which contains the acidic composition discussed above and an additive. Other aspects of the present invention pertain to a prepared nutriment containing a nutriment material and absorbed therein or adsorbed thereon is the acidic composition discussed above or the adduct discussed above. Another aspect of the present invention involves method to reduce biological contaminants in a nutriment material.

DETAILED DESCRIPTION

One embodiment of the present invention involves a highly acidic metalated mixture of inorganic acids ("HAMMIA"). The composition has an acidic pH, and can be isolated from a mixture prepared by mixing ingredients comprising a salt of phosphoric acid, and a preformed, or in-situ generated, solution or suspension of an acidic sparingly-soluble Group IIA complex ("AGIIS"), wherein the solution or suspension of AGIIS is in an amount sufficient to render the acidic pH of the composition to be less than about 2. Another embodiment of the present invention involves a composition having an acidic pH, the composition is isolated from a mixture prepared by mixing ingredients comprising a salt of phosphoric acid, and a preformed, or in-situ generated, solution or suspension of AGIIS, wherein the solution or suspension of AGIIS is in an amount in excess of the amount required to completely convert the salt of phosphoric acid to phosphoric acid. Still another embodiment of the present invention involves an adduct which contains an additive and the acidic composition of the present invention. Other aspects of the present invention pertain to a prepared nutriment containing a nutriment material and absorbed therein or adsorbed thereon is the acidic composition or the adduct of the present invention. Another aspect of the present invention involves method to reduce biological contaminants in a nutriment material.

The acidic, or low pH, solution of sparingly-soluble Group IIA complexes ("AGIIS") may have a suspension of very fine particles and the term "low pH" means the pH is below 7, in the acidic region. The AGIIS has a certain acid normality but does not have the same dehydrating behavior as a saturated calcium sulfate in sulfuric acid having the same normality. In other words, the AGIIS has a certain acid normality but does not char sucrose as readily as does a saturated solution of calcium sulfate in sulfuric acid having the same normality. Further, the AGIIS has low volatility at room temperature and pressure. It is less corrosive to a human skin than sulfuric acid saturated with

calcium sulfate having the same acid normality. Not intending to be bound by the theory, it is believed that one embodiment of AGIIS comprises near-saturated, saturated, or super-saturated calcium, sulfate anions or variations thereof, and/or complex ions containing calcium, sulfates, and/or variations thereof.

The term "complex," as used herein, denotes a composition wherein individual constituents are associated. "Associated" means constituents are bound to one another either covalently or non-covalently, the latter as a result of hydrogen bonding or other inter-molecular forces. The constituents may be present in ionic, non-ionic, hydrated or other forms.

The AGIIS can be prepared in several ways. Some of the methods involve the use of Group IA hydroxide but some of syntheses are devoid of the use of any added Group IA hydroxide, although it is possible that a small amount of Group IA metal may be present as "impurities." The preferred way of manufacturing AGIIS is not to add Group IA hydroxide to the mixture. As the phrase implies, AGIIS is highly acidic, ionic, with a pH of below about 7, preferably below about 2.

A preferred method of preparing AGIIS involves mixing a mineral acid with a Group IIA hydroxide, or with a Group IIA salt of a dibasic acid, or with a mixture of the two Group IIA materials. In the mixing, a salt of Group IIA is also formed. Preferably, the starting Group IIA material or materials selected will give rise to, and form, the Group IIA salt or salts that are sparingly soluble in water. The preferred mineral acid is sulfuric acid, the preferred Group IIA hydroxide is calcium hydroxide, and the preferred Group IIA salt of a dibasic acid is calcium sulfate. Other examples of Group IIA salt include calcium oxide, calcium carbonate, and "calcium bicarbonate."

Thus, for example, AGIIS can be prepared by mixing or blending starting materials given in one of the following scheme with good reproducibility:

- (1) H_2SO_4 and $\text{Ca}(\text{OH})_2$;
- 5 (2) H_2SO_4 , $\text{Ca}(\text{OH})_2$, and CaCO_3 ;
- (3) H_2SO_4 , $\text{Ca}(\text{OH})_2$, CaCO_3 , and CO_2 (gas);
- (4) H_2SO_4 , CaCO_3 , and $\text{Ca}(\text{OH})_2$;
- (5) H_2SO_4 , $\text{Ca}(\text{OH})_2$, and CaSO_4 ;
- (6) H_2SO_4 , CaSO_4 , CaCO_3 , and $\text{Ca}(\text{OH})_2$;
- 10 (7) H_2SO_4 , CaSO_4 , CaCO_3 , and CO_2 (gas); and
- (8) H_2SO_4 , CaSO_4 , CaCO_3 , CO_2 (gas), and $\text{Ca}(\text{OH})_2$.

Preferably, AGIIS is prepared by mixing calcium hydroxide with concentrated sulfuric acid, with or without an optional Group IIA salt of a dibasic acid (such as calcium sulfate) added to the sulfuric acid. The optional calcium sulfate can be added to the concentrated sulfuric acid prior to the introduction of calcium hydroxide into the blending mixture. The addition of calcium sulfate to the concentrated sulfuric acid appears to reduce the amount of calcium hydroxide needed for the preparation of AGIIS. Other optional reactants include calcium carbonate and gaseous carbon dioxide being bubbled into the mixture. Regardless of the use of any optional reactants, it was found that the use of calcium hydroxide is desirable.

One preferred method of preparing AGIIS can be described briefly as: Concentrated sulfuric acid is added to chilled water (8° - 12°C) in the reaction vessel, then, with stirring, calcium sulfate is added to the acid in chilled water to give a mixture. Temperature control is paramount to this process. To this stirring mixture is then added a slurry of calcium hydroxide in water. The solid formed from the mixture is then removed. This method involves the use of sulfuric acid, calcium sulfate, and calcium hydroxide, and it has several unexpected advantages. Firstly, this reaction is not violent and is not exceedingly exothermic. Besides being easy to control and easy to reproduce, this reaction uses ingredients each of which has been reviewed by the U.S. Food and Drug Administration ("U.S. FDA") and determined to be "generally recognized as safe" ("GRAS"). As such, each of these ingredients can be added directly to food, subject, of course, to certain limitations. Under proper concentration, each of these ingredients can be used as processing aids and in food contact applications. Their use is limited only by product suitability and Good Manufacturing Practices ("GMP"). The AGIIS so prepared is thus safe for animal consumption, safe for processing aids, and safe in food contact applications. Further, the AGIIS reduces biological contaminants in not only inhibiting the growth of, and killing, microorganisms but also destroying the toxins formed and generated by the microorganisms. The AGIIS formed can also preserve, or extend the shelf-life of, consumable products, be they plant, animal, pharmaceutical, or biological products. It also preserves or improves the organoleptic quality of a beverage, a plant product or an animal product. It also possesses certain healing and therapeutic properties.

The sulfuric acid used is usually 95-98% FCC Grade (about 35-37 *N*). The amount of concentrated sulfuric acid can range from about 0.05 *M* to about 18 *M* (about 0.1 *N* to about 36 *N*), preferably from about 1 *M* to about 5 *M*. It is application specific. The term "*M*" used denotes molar or moles per liter.

Normally, a slurry of finely ground calcium hydroxide suspended in water (about 50% of w/v) is the preferred way of introducing the calcium hydroxide, in increments, into the a stirring solution of sulfuric acid, with or without the presence of calcium sulfate. Ordinarily, the reaction is carried out
5 below 40°C, preferably below room temperature, and more preferably below 10°C. The time to add calcium hydroxide can range from about 1 hour to about 4 hours. The agitation speed can vary from about 600 to about 700 rpm or higher. After the mixing, the mixture is filtered through a 5 micron filter. The filtrate is then allowed to sit overnight and the fine sediment is removed by
10 decantation.

The calcium hydroxide used is usually FCC Grade of about 98% purity. For every mole of concentrated acid, such as sulfuric acid, the amount, in mole, of calcium hydroxide used is application specific and ranges from about 0.1 to about 1.

15 The phosphoric acid used is usually from JT Baker of about 85-88%.

The calcium monohydrogen phosphate is usually of 98-99%; and the calcium phosphate (“the tribasic”) is obtained from Mallinckrodt. Other phosphate salts used are all of reagent grade.

20 The optional calcium carbonate is normally FCC Grade having a purity of about 98%. When used with calcium hydroxide as described above, for every mole of concentrated acid, such as sulfuric acid, the amount, in mole, of calcium carbonate ranges from about 0.001 to about 0.2, depending on the amount of calcium hydroxide used.

25 The optional carbon dioxide is usually bubbled into the slurry containing calcium hydroxide at a speed of from about 1 to about 3 pounds pressure. The carbon dioxide is bubbled into the slurry for a period of from about 1 to about 3

hours. The slurry is then added to the reaction vessel containing the concentrated sulfuric acid.

5 Another optional ingredient is calcium sulfate, a Group IIA salt of a dibasic acid. Normally, dihydrated calcium sulfate is used. As used in this application, the phrase “calcium sulfate,” or the formula “CaSO₄,” means either anhydrous or hydrated calcium sulfate. The purity of calcium sulfate (dihydrate) used is usually 95-98% FCC Grade. The amount of calcium sulfate, in moles per liter of concentrated sulfuric acid ranges from about 0.005 to about 0.15, preferably from about 0.007 to about 0.07, and more preferably from about 0.007 to about 0.04. It is application specific.

In the event that CaSO₄ is used for the reaction by adding it to the solution of concentrated H₂SO₄, the amount of CaSO₄, in grams per liter of solution based on final volume, has the following relationship:

	<u>Final AGIIS Acid Normality <i>N</i></u>	<u>Amount of CaSO₄ in g/l</u>
15	1 - 5	5
	6-10	4
	11-15	3
	16-20	2
	21-36	1

20 The AGIIS obtained could have an acid normality range of from about 0.05 to about 31; the pH of lower than 0; boiling point of from about 100 to about 106°C; freezing point of from about -8°C to about 0°C.

AGIIS obtained from using the reaction of H₂SO₄/Ca(OH)₂/CaSO₄ had the following analyses (average):

AGIIS With Final Acid Normality of 1.2 N, pH of -0.08

H_3O^+ , 2.22%; Ca, 602 ppm; SO_4 , 73560 ppm; K, 1.36 ppb; impurities of 19.68 ppm, and neither Na nor Mg was detected.

AGIIS With Final Acid Normality of about 29 N, pH of about -1.46

- 5 H_3O^+ , 30.68%; Ca, 52.9 ppm; SO_4 , 7356000 ppm; K, 38.02 ppb; and neither Na nor Mg was detected.

10 Aqueous solutions of other alkalis or bases, such as Group IA hydroxide solution or slurry and Group IIA hydroxide solution or slurry can be used. Groups IA and IIA refer to the two Groups in the periodical table. The use of Group IIA hydroxide is preferred. Preferably, the salts formed from using Group IIA hydroxides in the reaction are sparingly soluble in water. It is also preferable to use only Group IIA hydroxide as the base without the addition of Group IA hydroxide.

15 After the reaction, the resultant concentrated acidic solution with a relatively low pH value, typically below pH 1, can then be diluted with de-ionized water to the desired pH value, such as pH of about 1 or about 1.8.

20 As discussed above, AGIIS has relatively less dehydrating properties (such as charring sucrose) as compared to the saturated solution of CaSO_4 in the same concentration of H_2SO_4 . Further, the stability and non-corrosive nature of the AGIIS of the present invention can be illustrated by the fact that a person can put his or her hand into this solution with a pH of less than 0.5 and, yet, his or her hand suffers no irritation, and no injury. If, on the other hand, one places his or her hand into a solution of sulfuric acid of pH of less than 0.5, an irritation would occur within a relatively short span of time. A solution of 28 N
25 of sulfuric acid saturated with calcium sulfate will cause chemical burn to a human skin after a few seconds of contact. In contrast, AGIIS solution of the

same normality would not cause chemical burn to a human skin even after in contact for 5 minutes. The AGIIS does not seem to be corrosive when being brought in contact with the environmental protective covering of plants (cuticle) and animals (skin). AGIIS has low volatility at room temperature and pressure. Even as concentrated as 29 *N*, the AGIIS has no odor, does not give off fumes in the air, and is not irritating to a human nose when one smells this concentrated solution.

The “additive” of the present invention appears to enhance, and also appears to be synergistic to, the effectiveness of the acidic composition of the present invention. Examples of the additive include alcohol, organic acid, periodic acid, and surfactant. The amount of additive added to the AGIIS varies depending on the desired final weight percent of the additive in the final adduct composition. The weight percent of additive needed for the adduct composition of the present invention can vary from about 0.01 to about 99.99, based on the total weight of the final adduct composition. The alcohol additive preferred for the present invention includes methanol, ethanol, 1-propanol, 2-propanol, and other lower alkyl alcohols.

Organic acid additive of the present invention includes carboxylic acid. A carboxylic acid is an organic compound containing the -COOH group, i.e., a carbonyl attached to a hydroxyl group. Preferred organic acids for the present invention include lactic acid, acetic acid, propionic acid, oxalic acid, sorbic acid, butyric acid, benzoic acid, glycolic acid, peracetic acid, and a mixture thereof.

A surfactant for the present invention is a surface-active agent. It is usually an organic compound consisting of two parts: One, a hydrophobic portion, usually including a long hydrocarbon chain; and two, a hydrophilic portion which renders the compound sufficiently soluble or dispersible in water or another polar solvent. Surfactants are usually classified into: (1) anionic,

where the hydrophilic moiety of the molecule carries a negative charge; (2) cationic, where this moiety of the molecule carries a positive charge; and (3) non-ionic, which do not dissociate, but commonly derive their hydrophilic moiety from polyhydroxy or polyethoxy structures. Other surfactants include ampholytic and zwitterionic surfactants. A preferred surfactant for the present invention includes polysorbates (Tween 80).

Unless otherwise defined, the amount of each ingredient or component of the present invention is based on the weight percent of the final composition, usually the concentrate before further dilution to achieve the desired pH of about 1.8. The AGIIS having a pH of about 1.8 is usually further diluted with water before applying to an animal product or a plant product.

A "biological contaminant" is a biologically substance, such as a biological organism of the product of a biological organism, that contaminates the environment and useful product, it is a biological material of an extraneous nature. This biological contaminant results in making the environment or product hazardous. A biological contaminant includes a microbe, a mold, and other infectious matter. A microbe is a very minute organism, and both microscopic and ultramicroscopic organisms, including spirochetes, bacteria, rickettsiae, and viruses.

The composition of the present invention was found to be a "preservative." The composition is not corrosive; however, it can create an environment where destructive micro-organisms cannot live and propagate. The utility of this method of preservation is that additional chemicals do not have to be added to the food or other substance to be preserved because the inherent low pH of the mixture is preservative. Since preservative chemicals do not have to be added to the food substance, taste is improved and residues are avoided. Organoleptic testing of a number of freshly preserved and previously preserved food stuffs have revealed the addition of composition improves taste

and eliminates preservative flavors. The term "organoleptic" means making an impression based upon senses of an organ or the whole organism.

The following examples are provided to further illustrate this invention and the manner in which it may be carried out. It will be understood, however, that the specific details given in the examples have been chosen for purposes of illustration only and not be construed as limiting the invention. Unless otherwise defined, the amount of each ingredient or component of the present invention is based on the weight percent of the final composition.

Example 1

AGIIS Having An Acid Normality of 1.2 To 1.5 Prepared By The Method Of $\text{H}_2\text{SO}_4/\text{Ca}(\text{OH})_2$.

An amount of 1055 ml (19.2 moles, after purity adjustment and taking into account the amount of acid neutralized by base) of concentrated sulfuric acid (FCC Grade, 95-98% purity) was slowly added with stirring, to 16.868 L of RO/DI water in each of reaction flasks a, b, c, e, and f. The amount of water had been adjusted to allow for the volume of acid and the calcium hydroxide slurry. The mixture in each flask was mixed thoroughly. Each of the reaction flasks was chilled in an ice bath and the temperature of the mixture in the reaction flask was about 8-12°C. The mixture was continuously stirred at a rate of about 700 rpm.

Separately, a slurry was made by adding RO/DI water to 4 kg of calcium hydroxide (FCC Grade, 98% purity) making a final volume of 8 L. The mole ratio of calcium hydroxide to concentrated sulfuric acid was determined to be 0.45 to 1. The slurry was a 50% (w/v) mixture of calcium hydroxide in water. The slurry was mixed well with a high-shear-force mixer until the slurry

appeared uniform. The slurry was then chilled to about 8-12°C in an ice bath and continuously stirred at about 700 rpm.

5 To each of the reaction flasks was added 150 ml of the calcium hydroxide slurry every 20 minutes until 1.276 L (i.e. 638 g dry weight, 8.61 moles, of calcium hydroxide) of the slurry had been added to each reaction vessel. The addition was again accompanied by efficient mixing at about 700 rpm.

10 After the completion of the addition of the calcium hydroxide to the reaction mixture in each reaction vessel, the mixture was filtered through a 5-micron filter.

The filtrate was allowed to sit for 12 hours, the clear solution was decanted to discard any precipitate formed. The resulting product was AGIIS having an acid normality of 1.2-1.5.

Example 2

15 AGIIS Having An Acid Normality Of 2 Prepared By The Method Of $\text{H}_2\text{SO}_4/\text{Ca}(\text{OH})_2/\text{CaSO}_4$.

20 For the preparation of 1 L of 2 N AGIIS, an amount of 79.5 ml (1.44 moles, after purity adjustment and taking into account the amount of acid to be neutralized by base) of concentrated sulfuric acid (FCC Grade, 95-98% purity) was slowly added, with stirring, to 854 ml of RO/DI water in a 2 L reaction flask. Five grams of calcium sulfate (FCC Grade, 95% purity) was then added slowly and with stirring to the reaction flask. The mixture was mixed thoroughly. At this point, analysis of the mixture would usually indicate an acid normality of 2.88. The reaction flask was chilled in an ice bath and the

temperature of the mixture in the reaction flask was about 8-12°C. The mixture was continuously stirred at a rate of about 700 rpm.

5 Separately, a slurry was made by adding 50 ml of RO/DI water to 33.26 g (0.44 mole, after purity adjustment) of calcium hydroxide (FCC Grace, 98% purity) making a final volume of 66.53 ml. The mole ratio of calcium hydroxide to concentrated sulfuric acid was determined to be 0.44 to 1. The slurry was mixed well with a high-shear-force mixer until the slurry appeared uniform. The slurry was then chilled to about 8-12°C in an ice bath and continuous stirred at about 700 rpm.

10 The slurry was then slowly added over a period of 2-3 hours to the mixture, still chilled in an ice bath and being stirred at about 700 rpm.

15 After the completion of the addition of slurry to the mixture, the product was filtered through a 5-micron filter. It was normal to observe a 20% loss in volume of the mixture due to the retention of the solution by the salt and removal of the salt.

The filtrate was allowed to sit for 12 hours, and the clear solution was then decanted to discard any precipitate formed. The resulting product was AGIIS having an acid normality of 2.

Example 3

20 AGIIS Having An Acid Normality Of 12 Prepared By The Method Of $\text{H}_2\text{SO}_4/\text{Ca}(\text{OH})_2/\text{CaSO}_4$.

For the preparation of 1 L of 12 N AGIIS, an amount of 434 ml (7.86 moles, after purity adjustment and taking into account amount of acid neutralized by base) of concentrated sulfuric acid (FCC Grade, 95-98% purity)

was slowly added, with stirring, to 284.60 ml of RO/DI water in a 2 L reaction flask. Three grams of calcium sulfate (FCC Grade, 95% purity) was then added slowly and with stirring to the reaction flask. The mixture was mixed thoroughly. The reaction flask was chilled in an ice bath and the temperature of the mixture in the reaction flask was about 8-12°C. The mixture was continuously stirred at a rate of about 700 rpm.

Separately, a slurry was made by adding 211 ml of RO/DI water to 140.61 g (1.86 moles, after purity adjustment) of calcium hydroxide (FCC Grade, 98% purity) making a final volume of 281.23 ml. The mole ratio of calcium hydroxide to concentrated sulfuric acid was determined to be 0.31. The slurry was mixed well with a high-shear-force mixer until the slurry appeared uniform. The slurry was then chilled to about 8-12°C in an ice bath and continuous stirred at about 700 rpm.

The slurry was then slowly added over a period of 2-3 hours to the acid mixture, still chilled in an ice bath and being stirred at about 700 rpm.

After the completion of the addition of slurry to the mixture, the product was filtered through a 5-micron filter. It was normal to observe a 20% loss in volume of the mixture due to the retention of the solution by the salt and removal of the salt.

The filtrate was allowed to sit for 12 hours, and the clear solution was then decanted to discard any precipitate formed. The resulting product was AGIIS having an acid normality of 12.

Example 4

General Procedure 1. Formation Of A Phosphoric Acid HAMMIA Using Pre-Formed AGIIS.

5 The phosphate salt of a divalent metal chosen from List A below (1.00 mole equivalents) is suspended in sufficient deionized water to make a final volume of 625 mL per mole of phosphate ions. The mixture may be sonicated or heated as necessary to aid solubilization of the sparingly soluble phosphate salt. To this stirred suspension, a solution of AGIIS containing the desired concentration of acid (3.05 moles of hydrogen ion per mole of phosphate ion; 2.05 moles of hydrogen ion per mole of hydrogen phosphate ion; 1.05 moles of hydrogen ion per mole of dihydrogen phosphate ion) is added in 10-mL aliquots with the pH being monitored after each addition. Copious precipitates of calcium sulfate form beginning at pH 2. The addition of AGIIS solution may be discontinued as soon as the desired pH is reached. After the addition of the acid is complete, the mixture is stirred for one hour. The agitation is then stopped and the mixture is allowed to settle overnight (approximately 18 hours). The suspended solids are removed by centrifugation at 16000 rpm for 30 minutes. 10 The supernatant solution is the HAMMIA.

List A: Phosphate Salts

Mg₃(PO₄)₂, MgHPO₄, Mg(H₂PO₄)₂

Ca₃(PO₄)₂, CaHPO₄, Ca(H₂PO₄)₂

Mn₃(PO₄)₂, MnHPO₄, Mn(H₂PO₄)₂

20 Fe₃(PO₄)₂, FeHPO₄, Fe(H₂PO₄)₂

Co₃(PO₄)₂, CoHPO₄, Co(H₂PO₄)₂

Ni₃(PO₄)₂, NiHPO₄, Ni(H₂PO₄)₂

Cu₃(PO₄)₂, CuHPO₄, Cu(H₂PO₄)₂

Zn₃(PO₄)₂, ZnHPO₄, Zn(H₂PO₄)₂

25

Example 5

General Procedure 2. Formation Of A Phosphoric Acid HAMMIA Using AGIIS Formed In Situ.

5 A mixture of calcium hydroxide (1.00 mole equivalents) and the
 phosphate salt of a divalent metal chosen from List A below (1.00 mole
 equivalents) is suspended in sufficient deionized water to make a final volume
 of approximately 400 mL per mole of metal ions. The mixture may be
 sonicated or heated as necessary to aid solubilization of the sparingly soluble
 metal salts. To this stirred suspension, concentrated sulfuric acid (5.05 mole
 equivalents of hydrogen ion per mole of phosphate ion) is added in 10-mL
 aliquots with the pH being monitored after each addition. The addition of acid
 may be discontinued when the desired pH is reached. After the addition of the
 10 acid is complete, the mixture is stirred for one hour. The agitation is then
 stopped and the mixture is allowed to settle overnight (approximately 18 hours).
 The suspended solids are removed by centrifugation at 16000 rpm for 30
 minutes. The supernatant solution is the HAMMIA.

List A: Phosphate Salts

15 $Mg_3(PO_4)_2$, $MgHPO_4$, $Mg(H_2PO_4)_2$
 $Ca_3(PO_4)_2$, $CaHPO_4$, $Ca(H_2PO_4)_2$
 $Mn_3(PO_4)_2$, $MnHPO_4$, $Mn(H_2PO_4)_2$
 $Fe_3(PO_4)_2$, $FeHPO_4$, $Fe(H_2PO_4)_2$
 $Co_3(PO_4)_2$, $CoHPO_4$, $Co(H_2PO_4)_2$
 20 $Ni_3(PO_4)_2$, $NiHPO_4$, $Ni(H_2PO_4)_2$
 $Cu_3(PO_4)_2$, $CuHPO_4$, $Cu(H_2PO_4)_2$
 $Zn_3(PO_4)_2$, $ZnHPO_4$, $Zn(H_2PO_4)_2$

Example 6

25 General Procedure 3. Formation Of A Phosphoric Acid HAMMIA
 Containing A Monovalent Metal Using Pre-formed AGIIS.

The phosphate salt of a divalent metal chosen from List A below (1.00
 mole equivalents) and the phosphate salt of a monovalent metal chosen from

List B below (≤ 1.00 mole equivalents) is suspended in sufficient deionized water to make a final volume of 625 mL per mole of phosphate ions. The mixture may be sonicated or heated as necessary to aid solubilization of the sparingly soluble divalent metal phosphate salt. To this stirred suspension, a solution of AGIIS containing the desired concentration of acid (3.05 moles of hydrogen ion per mole of phosphate ion; 2.05 moles of hydrogen ion per mole of hydrogen phosphate ion; 1.05 moles of hydrogen ion per mole of dihydrogen phosphate ion) is added in 10-mL aliquots with the pH being monitored after each addition. Copious precipitates of calcium sulfate form beginning at pH 2. The addition of AGIIS solution may be discontinued as soon as the desired pH is reached. After the addition of the acid is complete, the mixture is stirred for one hour. The agitation is then stopped and the mixture is allowed to settle overnight (approximately 18 hours). The suspended solids are removed by centrifugation at 16000 rpm for 30 minutes. The supernatant solution is the HAMMIA.

List A:
Divalent Metal Phosphate Salts

$Mg_3(PO_4)_2$, $MgHPO_4$, $Mg(H_2PO_4)_2$
 $Ca_3(PO_4)_2$, $CaHPO_4$, $Ca(H_2PO_4)_2$
 $Mn_3(PO_4)_2$, $MnHPO_4$, $Mn(H_2PO_4)_2$
 $Fe_3(PO_4)_2$, $FeHPO_4$, $Fe(H_2PO_4)_2$
 $Co_3(PO_4)_2$, $CoHPO_4$, $Co(H_2PO_4)_2$
 $Ni_3(PO_4)_2$, $NiHPO_4$, $Ni(H_2PO_4)_2$
 $Cu_3(PO_4)_2$, $CuHPO_4$, $Cu(H_2PO_4)_2$
 $Zn_3(PO_4)_2$, $ZnHPO_4$, $Zn(H_2PO_4)_2$

List B:
Monovalent Metal Phosphate Salts

Li_3PO_4 , Li_2HPO_4 , LiH_2PO_4
 Na_3PO_4 , Na_2HPO_4 , NaH_2PO_4
 K_3PO_4 , K_2HPO_4 , KH_2PO_4

Example 7

General Procedure 4. Formation Of A Phosphoric Acid HAMMIA Containing A Monovalent Metal Using AGIIS Formed In Situ.

5 A mixture of calcium hydroxide (1.00 mole equivalents) and the phosphate salt of a divalent metal chosen from List A below (1.00 mole equivalents) is suspended in sufficient deionized water to make a final volume of approximately 400 mL per mole of metal ions. The phosphate salt of a monovalent metal chosen from List B below (≤ 1.00 mole equivalents) is added to the mixture. The mixture may be sonicated or heated as necessary to aid solubilization of the sparingly soluble divalent metal salts. To this stirred suspension, concentrated sulfuric acid (5.05 mole equivalents of hydrogen ion per mole of phosphate ion) is added in 10-mL aliquots with the pH being monitored after each addition. The addition of acid may be discontinued when the desired pH is reached. After the addition of the acid is complete, the mixture is stirred for one hour. The agitation is then stopped and the mixture is allowed to settle overnight (approximately 18 hours). The suspended solids are removed by centrifugation at 16000 rpm for 30 minutes. The supernatant solution is the HAMMIA.

List A:
Divalent Metal Phosphate Salts

List B:
Monovalent Metal Phosphate Salts

Mg ₃ (PO ₄) ₂ , MgHPO ₄ , Mg(H ₂ PO ₄) ₂	Li ₃ PO ₄ , Li ₂ HPO ₄ , LiH ₂ PO ₄
Ca ₃ (PO ₄) ₂ , CaHPO ₄ , Ca(H ₂ PO ₄) ₂	Na ₃ PO ₄ , Na ₂ HPO ₄ , NaH ₂ PO ₄
Mn ₃ (PO ₄) ₂ , MnHPO ₄ , Mn(H ₂ PO ₄) ₂	K ₃ PO ₄ , K ₂ HPO ₄ , KH ₂ PO ₄
Fe ₃ (PO ₄) ₂ , FeHPO ₄ , Fe(H ₂ PO ₄) ₂	
Co ₃ (PO ₄) ₂ , CoHPO ₄ , Co(H ₂ PO ₄) ₂	
Ni ₃ (PO ₄) ₂ , NiHPO ₄ , Ni(H ₂ PO ₄) ₂	
Cu ₃ (PO ₄) ₂ , CuHPO ₄ , Cu(H ₂ PO ₄) ₂	
Zn ₃ (PO ₄) ₂ , ZnHPO ₄ , Zn(H ₂ PO ₄) ₂	

Example 8

General Procedure 5. Formation Of A Phosphoric Acid HAMMIA Containing A Monovalent Metal And An Additive Acid Using Pre-formed AGIIS.

- 5 One or more of the acids from List C below (up to 6 mole equivalents), the phosphate salt of a divalent metal chosen from List A below (1.00 mole equivalents) and the phosphate salt of a monovalent metal chosen from List B below (≤ 1.00 mole equivalents) are suspended in sufficient deionized water to make a final volume of 625 mL per mole of phosphate ions. The mixture may
- 10 be sonicated or heated as necessary to aid solubilization of the sparingly soluble divalent metal phosphate salt. To this stirred suspension, a solution of AGIIS containing the desired concentration of acid (3.05 moles of hydrogen ion per mole of phosphate ion; 2.05 moles of hydrogen ion per mole of hydrgen phosphate ion; 1.05 moles of hydrogen ion per mole of dihydrogen phosphate
- 15 ion) is added in 10-mL aliquots with the pH being monitored after each addition. Copious precipitates of calcium sulfate form beginning at pH 2. The addition of AGIIS solution may be discontinued as soon as the desired pH is reached. After the addition of the acid is complete, the mixture is stirred for one

hour. The agitation is then stopped and the mixture is allowed to settle overnight (approximately 18 hours). The suspended solids are removed by centrifugation at 16000 rpm for 30 minutes. The supernatant solution is the HAMMIA.

List A:
Divalent Metal Phosphate Salts

Mg₃(PO₄)₂, MgHPO₄, Mg(H₂PO₄)₂
 Ca₃(PO₄)₂, CaHPO₄, Ca(H₂PO₄)₂
 Mn₃(PO₄)₂, MnHPO₄, Mn(H₂PO₄)₂
 Fe₃(PO₄)₂, FeHPO₄, Fe(H₂PO₄)₂
 Co₃(PO₄)₂, CoHPO₄, Co(H₂PO₄)₂
 Ni₃(PO₄)₂, NiHPO₄, Ni(H₂PO₄)₂
 Cu₃(PO₄)₂, CuHPO₄, Cu(H₂PO₄)₂
 Zn₃(PO₄)₂, ZnHPO₄, Zn(H₂PO₄)₂

List B:
Monovalent Metal Phosphate Salts

Li₃PO₄, Li₂HPO₄, LiH₂PO₄
 Na₃PO₄, Na₂HPO₄, NaH₂PO₄
 K₃PO₄, K₂HPO₄, KH₂PO₄

List C:
Additive Acids

formic acid, acetic acid, propionic
 acid, butyric acid, malic acid,
 glycolic acid, maleic acid, gluconic
 acid, periodic acid, peracetic acid,
 monoperphthalic acid, benzoic acid,
 sorbic acid, oxalic acid.

Example 9

General Procedure 6. Formation Of A Phosphoric Acid HAMMIA
 Containing A Monovalent Metal And An Additive Acid Using AGIIS Formed
 In Situ.

A mixture of calcium hydroxide (1.00 mole equivalents) and the
 phosphate salt of a divalent metal chosen from List A below (1.00 mole
 equivalents) is suspended in sufficient deionized water to make a final volume
 of approximately 400 mL per mole of metal ions. One or more of the acids

from List C below (up to 6 mole equivalents), and phosphate salt of a monovalent metal chosen from List B below (≤ 1.00 mole equivalents) is added to the mixture. The mixture may be sonicated or heated as necessary to aid solubilization of the sparingly soluble divalent metal salts. To this stirred suspension, concentrated sulfuric acid (5.05 mole equivalents of hydrogen ion per mole of phosphate ion) is added in 10-mL aliquots with the pH being monitored after each addition. The addition of acid may be discontinued when the desired pH is reached. After the addition of the acid is complete, the mixture is stirred for one hour. The agitation is then stopped and the mixture is allowed to settle overnight (approximately 18 hours). The suspended solids are removed by centrifugation at 16000 rpm for 30 minutes. The supernatant solution is the HAMMIA.

List A:

Divalent Metal Phosphate Salts

$\text{Mg}_3(\text{PO}_4)_2$, MgHPO_4 , $\text{Mg}(\text{H}_2\text{PO}_4)_2$

$\text{Ca}_3(\text{PO}_4)_2$, CaHPO_4 , $\text{Ca}(\text{H}_2\text{PO}_4)_2$

$\text{Mn}_3(\text{PO}_4)_2$, MnHPO_4 , $\text{Mn}(\text{H}_2\text{PO}_4)_2$

$\text{Fe}_3(\text{PO}_4)_2$, FeHPO_4 , $\text{Fe}(\text{H}_2\text{PO}_4)_2$

$\text{Co}_3(\text{PO}_4)_2$, CoHPO_4 , $\text{Co}(\text{H}_2\text{PO}_4)_2$

$\text{Ni}_3(\text{PO}_4)_2$, NiHPO_4 , $\text{Ni}(\text{H}_2\text{PO}_4)_2$

$\text{Cu}_3(\text{PO}_4)_2$, CuHPO_4 , $\text{Cu}(\text{H}_2\text{PO}_4)_2$

$\text{Zn}_3(\text{PO}_4)_2$, ZnHPO_4 , $\text{Zn}(\text{H}_2\text{PO}_4)_2$

List B:

Monovalent Metal Phosphate Salts

Li_3PO_4 , Li_2HPO_4 , LiH_2PO_4

Na_3PO_4 , Na_2HPO_4 , NaH_2PO_4

K_3PO_4 , K_2HPO_4 , KH_2PO_4

List C:

Additive Acids

formic acid, acetic acid, propionic acid, butyric acid, malic acid, glycolic acid, maleic acid, gluconic acid, periodic acid, peracetic acid, monoperphthalic acid, benzoic acid, sorbic acid, oxalic acid.

Example 10

Preparation Of HAMMIA and Adduct Containing HAMMIA And Additive

Procedure A

5 The phosphate salt of a divalent metal chosen from List C below (1.00 mole equivalents) is suspended in sufficient deionized water to make a final volume of 625 mL per mole of phosphate ions. The mixture may be sonicated 30 minutes or heated as necessary to aid solubilization of the sparingly soluble phosphate salt. To this stirred suspension concentrated sulfuric acid (167.5 mL per mole of phosphate ions, 97%, 3.05 mole equivalents) is added in 10-mL aliquots each 20 minutes. Below pH 2, copious precipitation of calcium sulfate begins. After the addition of the acid is complete, the mixture is stirred for one hour and the agitation is stopped and the mixture is allowed to settle overnight (approximately 18 hours). The suspended solids are removed by centrifugation at 16000 rpm for 30 minutes.

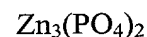
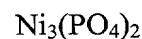
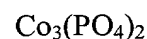
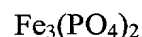
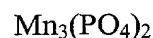
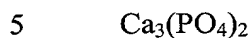
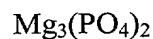
10

15

The supernatant resulting from this procedure using calcium phosphate, $\text{Ca}_3(\text{PO}_4)_2$, had a volume of approximately 1 L, a pH of approximately 0.0-0.5,

and contained approximately 1000 ppm Ca, 3.80×10^5 ppm SO_4 , and 1.14×10^5 ppm PO_4 .

List C: Phosphate Salts



Procedure B.

The monohydrogen phosphate salt of a divalent metal chosen from List D below (7.35 moles) is placed in an 8-L container and to deionized water (1.0 L) is added. The mixture is stirred using high shear force mixing during all subsequent additions. To this stirred suspension 1.45 L of a solution of AGIIS having an acid normality of 5.2 *N* is added in 10-mL aliquots. below pH 2, copious precipitation of calcium sulfate begins. After addition of 1.45 L of sulfuric acid, the pH of the mixture is approximately 1.0. After the addition of the acid is complete, a 2-L sample of the mixture is centrifuged at 15000 rpm for 20 minutes.

The supernatant resulting from this procedure using calcium monohydrogen phosphate (CaHPO_4) had a pH of approximately 1.23, and contained approximately 88 ppm Ca, 1800 ppm SO_4 , and 1.48×10^5 ppm PO_4 .

List D: Monohydrogen Phosphate Salts

MgHPO₄

CaHPO₄

MnHPO₄

5 FeHPO₄

CoHPO₄

NiHPO₄

CuHPO₄

ZnHPO₄

10 Procedure C.

The monohydrogen phosphate salt of a divalent metal chosen from List D above (11.0 moles) is placed in an 8-L container and deionized water (2.0 L) is added. The mixture is stirred using high shear force mixing during all subsequent additions. To this stirred suspension concentrated sulfuric acid (up to 500 mL, up to 9.15 moles) is added in 10-mL aliquots. The pH may be monitored, and the addition of sulfuric acid ceased when the desired pH is reached. The pH of the solution varies with the quantity of sulfuric acid added approximately as follows: pH 3.0, 40 mL; pH 2.0, 90 mL; pH 1.0, 240 mL; pH 0.5, 380 mL; pH 0.0 450 mL; pH<0, 470 mL. Below pH 2, copious precipitation of calcium sulfate occurs. After the addition of the sulfuric acid is complete, the mixture is centrifuged at 15000 rpm for 15-20 minutes.

The supernatant resulting from this procedure using calcium monohydrogen phosphate (CaHPO₄) and 500 mL of concentrated sulfuric acid had an acid concentration of approximately 7 N, and contained approximately 1.38x10⁴ ppm SO₄, 4.44x10⁵ ppm PO₄, 1.1 x 10³ ppm Ca.

Procedure D.

Concentrated phosphoric acid (1L, 16.8 moles) is placed in a container. The oxide, hydroxide, carbonate or basic carbonate salt of a divalent metal chosen from List E below (17.1 moles) is added in 50-g portions to the phosphoric acid and the mixture is well mixed after each addition. Water (2.9 L) is added as necessary to permit efficient mixing of the mixture. After the addition of the base and the water is complete, concentrated sulfuric acid (927 mL, 17.0 moles) is added to the stirred solution in 10-mL aliquots at a rate of 10 mL per 15 minutes. The pH of the solution varies with the quantity of sulfuric acid added approximately as follows: pH 3.0, 30 mL; pH 2.0, 120 mL; pH 1.0, 480 mL; pH 0.5, 640 mL; pH 0.0 710 mL; pH<0, 760 mL. Below pH 2, copious precipitation of calcium sulfate occurs. After the addition of the acid is complete, deionized water (500 mL) is added and the mixture is stirred well. Agitation is then stopped, and the mixture is allowed to settle overnight (approximately 18 hours). The suspended solids are removed by centrifugation at 15000 rpm for 20 minutes.

The supernatant resulting from this procedure using calcium hydroxide ($\text{Ca}(\text{OH})_2$) had a pH below 0.0, and contained approximately 250 ppm Ca, 1.00×10^5 ppm SO_4 , and 3.19×10^5 ppm PO_4 .

- 20 List E: Metal bases
MgO, $\text{Mg}(\text{OH})_2$, MgCO_3 , $x\text{MgO} \cdot y\text{MgCO}_3$
CaO, $\text{Ca}(\text{OH})_2$, CaCO_3
MnO, $\text{Mn}(\text{OH})_2$, MnCO_3 , $x\text{MnO} \cdot y\text{MnCO}_3$
FeO, $\text{Fe}(\text{OH})_2$, FeCO_3 , $x\text{FeO} \cdot y\text{FeCO}_3$
25 CoO, $\text{Co}(\text{OH})_2$, CoCO_3 , $x\text{CoO} \cdot y\text{CoCO}_3$
NiO, $\text{Ni}(\text{OH})_2$, NiCO_3 , $x\text{NiO} \cdot y\text{NiCO}_3$
CuO, $\text{Cu}(\text{OH})_2$, CuCO_3 , $x\text{CuO} \cdot y\text{CuCO}_3$
ZnO, $\text{Zn}(\text{OH})_2$, ZnCO_3 , $x\text{ZnO} \cdot y\text{ZnCO}_3$

Procedure D-1.

Propionic acid (110 mL, 1.48 mol) was dissolved in deionized water (890 mL) and a solution of AGIIS (5 N , 74 mL, 0.37 mol hydrogen ion) was added. This solution was stirred, and then solid calcium dihydrogen phosphate (25 g, 0.0214 moles) and calcium hydrogen phosphate (5 g, 0.184 moles) were added with vigorous stirring. As necessary, the mixture was centrifuged to remove suspended solids. The solution prepared by this method had a pH of approximately 1.5, and contained 2.6×10^4 ppm PO_4 , 3.1×10^3 ppm SO_4 , and 9.3×10^4 ppm $\text{C}_2\text{H}_5\text{CO}_2\text{H}$.

The same solution may be prepared as a five-fold concentrate by following the same procedure as modified below. The initial solution is prepared by mixing 550 mL (7.37 moles) of propionic acid and 450 mL of water. To this solution, AGIIS (5 N , 370 mL, 1.85 moles hydrogen ion) is added. This solution is stirred, and calcium dihydrogen phosphate (25 g, 0.107 moles) and calcium monohydrogen phosphate (125 g, 0.92 moles) are added portionwise with vigorous mixing. As necessary, suspended solids are removed from the final mixture by centrifugation. The resultant solution contains approximately 4.73×10^4 ppm PO_4 , 2.15×10^5 ppm SO_4 , and 4.11×10^5 ppm $\text{C}_2\text{H}_5\text{CO}_2\text{H}$. Dilution of this solution (200 mL) with deionized water (800 mL) gives a solution with a pH of approximately 1.1, and containing approximately 9.0×10^3 ppm PO_4 , 6.4×10^3 ppm SO_4 , and 7.6×10^4 ppm $\text{C}_2\text{H}_5\text{CO}_2\text{H}$.

Procedure D-2.

Propionic acid (110 mL, 1.48 mol) was dissolved in deionized water (890 mL) and a solution of AGIIS (5 N , 40 mL, 0.25 mol hydrogen ion) was added in 10-mL aliquots. This solution was stirred, and then solid sodium hydrogen phosphate (Na_2HPO_4 , 22 g, 0.155 moles) was added portionwise (4 x 5 g, 1 x 2 g) with vigorous stirring. After the addition of sodium hydrogen phosphate, an additional 45 mL of water was added to bring the total volume to

1.0 L. The solution prepared by this method had a pH of approximately 1.5, and contained 7.9×10^3 ppm PO_4 , 1.1×10^4 ppm SO_4 , and 1.0×10^5 ppm $\text{C}_2\text{H}_5\text{CO}_2\text{H}$.

The same solution may be prepared as a six-fold concentrate by following the same procedure as modified below. The initial solution is prepared by mixing 660 mL (8.84 moles) of propionic acid and 170 mL of water. To this solution, AGIIS (5 N, 240 mL, 1.2 moles hydrogen ion) is added. This solution is stirred, and sodium monohydrogen phosphate (Na_2HPO_4 , 132 g, 0.93 moles) is added portionwise with vigorous mixing.

Procedure D-3.

Propionic acid (110 mL, 1.48 mol) and lactic acid (100 mL, 85% in water, 103 g, 1.14 mol) were dissolved in deionized water (650 mL) and a solution of AGIIS (5 N, 28 mL, 0.14 mol hydrogen ion) was added in 10-mL aliquots. This solution is stirred, and then solid sodium hydrogen phosphate (71 g, 0.119 moles) was added portionwise with vigorous stirring. Water (90 mL) was added after addition of the sodium phosphate salts was complete. The solution prepared by this method contained approximately 6.5×10^3 ppm PO_4 , 7.2×10^3 ppm SO_4 , 1.0×10^5 ppm $\text{C}_2\text{H}_5\text{CO}_2\text{H}$ and 9.0×10^4 ppm $\text{CH}_3\text{CH}(\text{OH})\text{CO}_2\text{H}$.

The same solution may be prepared as a three-fold concentrate by following the same procedure as modified below. The initial solution is prepared by mixing 330 mL (4.03 moles) of propionic acid, 330 mL (308 g, 3.76 moles) of lactic acid, and 240 mL of water. A solution of AGIIS (5 N, 84 mL, 0.425 moles hydrogen ion) is added to the stirred solution. Solid sodium monohydrogen phosphate (52 g, 0.37 moles) is added portionwise with vigorous mixing. The resultant solution contains approximately 1.8×10^4 ppm PO_4 , 2.2×10^4 ppm SO_4 , and 3.6×10^5 ppm $\text{C}_2\text{H}_5\text{CO}_2\text{H}$ and 3.3×10^5 ppm

CH₃CH(OH)CO₂H. Dilution of this solution 1:3 with deionized water gives a solution containing approximately 5.8x10³ ppm PO₄, 7.0x10³ ppm SO₄, 1.0x10⁵ ppm C₂H₅CO₂H and 9.6x10⁴ ppm CH₃CH(OH)CO₂H.

5 The same three-fold concentrate may be prepared in gallon quantities by following the procedure as modified below. The initial solution is prepared by mixing 1250 mL of propionic acid, 1250 mL of 85% lactic acid, and 908 mL of water. A solution of AGIIS (5 N, 318 mL) is added to the stirred solution. Solid sodium monohydrogen phosphate (193 g) is added portionwise with vigorous mixing. Dilution of this solution 1:3 with deionized water gives a
10 solution with a pH of 1.5, and containing approximately 1.9x10³ ppm PO₄, 3.3x10³ ppm SO₄, 1.0x10⁵ ppm C₂H₅CO₂H and 1.1x10⁵ ppm CH₃CH(OH)CO₂H.

Procedure D-4

15 Calcium phosphate (500 g, 1.61 moles) was added to an 8-L container and a solution of AGIIS (1.0 L, 5 N, 5.0 moles hydrogen ion) was added dropwise at a rate of approximately 2 mL/minute. The mixture was stirred well, and deionized water (500 mL) was added to aid stirring. A further 500 mL of the AGIIS solution (5 N, 2.5 moles hydrogen ion) was added dropwise at a rate of approximately 2 mL/minute with vigorous stirring. The solids were removed
20 from the resultant mixture by centrifugation at 15000 rpm for 20 minutes. The supernatant solution was used as the HAMMIA.

The HAMMIA prepared by this method had a pH of 1.0-1.5, and contained approximately 1.2x10⁴ ppm Ca²⁺, 1.6x10³ ppm SO₄, and 1.5x10⁵ ppm PO₄.

Discussion

Although not wanting to be bound by any theory, the various adducts solutions containing HAMMIA and an additive acid as well as the various HAMMIA solutions were formed by the regeneration of phosphoric acid from its salts by a regenerating acid. The formation of the acidic solutions led to solutions that, when brought to a pH below 1.0, no longer had a substantial concentration of the metal ion (with calcium salts, the calcium ion concentration was around or below 1000 ppm, i.e. around or below 0.025 *M*). In most of the solutions prepared with calcium salts, the calcium ion concentration was below 200 ppm (0.005 *M*), when the pH was below 1. Thus, these with solutions pH<1 cannot, in general, be equated with a traditional buffer solution, where the concentrations of the metal salt are typically in the 0.1-0.5 *M* range. However, the anion concentrations are in the much higher concentration ranges; the simplest interpretation of the data would suggest that the solutions contain hydronium ion as the most prevalent cation present. Such solutions might meet the *pro forma* definition of a buffer, but such solutions will not behave as a functional buffer solution. It is noteworthy that the generation of these solutions appeared to pass through a buffer solution phase, where the addition of substantial volumes of the strong regenerating acid (usually AGIIS) had little effect on the pH of the mixture, but after the addition of the theoretical amount of regenerating acid, the pH dropped rapidly with additional regenerating acid. In the pH range above 1, and more especially above 1.5, the solution might act as a calcium dihydrogen phosphate buffer, and, as such, the calcium ion concentration may be much higher than in the pH < 1 solutions. Indeed, until the rapid drop of pH with added regenerating acid, it is possible to have quite high calcium ion concentrations of several thousand parts per million (as high as 0.3 *M*).

Example 11

Using Adduct Containing HAMMIA To Decontaminate Hot Dogs

In this experiment, the adduct solution containing HAMMIA, propionic acid, and lactic acid was prepared as described in Procedure D-3 for the "three-fold concentration." The acidic solution was diluted 1:3 with deionized water before use. The adduct solution contained lactate at approximately 100298 ppm, propionate at about 105314 ppm, sulfate at about 7960 ppm, and phosphate at about 7995 ppm. The pH of the adduct was 1.51.

The surface of each of a number of hot dogs was inoculated by dripping with 50 μ L of a *Listeria monocytogenes* culture containing 1.38×10^8 Colony Forming Unit ("CFU"), and then allowed to dry. In a treated group, the hot dogs were immersed in the solution of HAMMIA for 30 seconds. The excess solution was allowed to drain for 10 seconds. In a control group, the hot dogs were immersed in a saline solution and drained for the same length of time. Each group of hot dogs was packed into plastic a zip bag and stored at room temperature. After 24 hours, a comparison of the surviving *Listeria* colonies was made. The control group of hot dogs had an average of 3.5×10^9 CFU per hot dog, while the treated group had an average of 5.56×10^1 CFU per hot dog.

Example 12

Using Adduct Containing HAMMIA To Prolong The Shelf-Life Of Sausages And Hot Dogs.

In this experiment, the adduct solution containing HAMMIA was the same as in Example 11.

Three types of hot dogs (chicken, beef, and a beef, pork, chicken mixture) and beef sausage were used in this experiment. In a treated group, the four meat products were immersed in the adduct solution for 30 seconds, and then allowed to drain. A control group of meat products was used as purchased and was not immersed in any solution. Each group was packed into a plastic zip

bag and stored at room temperature. At 3 days and 6 days, the hot dogs and sausages were measured for total plaque count (“TPC”).

Microbial Presence on Untreated and Treated Meat Products at 3 and 6 days

<u>Meat product</u>	<u>Treated (CFU per meat product)</u>	<u>Untreated (CFU per meat product)</u>
<u>3 days:</u>		
*Chicken hot dog	$<1.1 \times 10^2$	2.08×10^9
Beef hot dog	$<1.1 \times 10^2$	4.5×10^8
Beef, pork, chicken hot dog	$<1.1 \times 10^2$	6.5×10^9
Sausage	$<1.1 \times 10^2$	4.11×10^5
<u>6 days:</u>		
Chicken hot dog	$<1.1 \times 10^2$	7.03×10^{10}
Beef hot dog	$<1.1 \times 10^2$	3.33×10^{10}
Beef, pork, chicken hot dog	$<1.1 \times 10^2$	5.92×10^{10}
Sausage	$<1.1 \times 10^2$	3.32×10^{10}

5 * 1.1×10^2 CFU per hot dog is considered undetectable in the study.

Example 13

Using Adduct Containing HAMMIA To Prevent The Growth Of Mold
On Pepperoni

10 In this experiment, the adduct solution containing HAMMIA was the
same as in Example 11.

Cooked pepperoni was cut into pieces seven inches in length. A treated group of pepperoni pieces was immersed in the adduct solution for 30 seconds and allowed to drain. A control group of pepperoni pieces was used as

5 purchased and was not immersed in any solution. All of the pepperoni pieces were packaged individually in plastic zip bags and stored at room temperature. After four days, both groups were examined visually. The control group of pepperoni pieces showed green mold growth on the surface, while the treated group did not. The treated group of pepperoni pieces did not show the growth of mold even after more than two months from the start of the experiment.

Example 14

Using Adduct Containing HAMMIA To Prevent Microbial Growth In Beef Steaks

10 In this experiment, the adduct solution containing HAMMIA was the same as in Example 11.

15 Beef steaks were injected with 1.16×10^3 CFU per 100 g of *E. coli* O157:H7 culture and 2.24×10^3 CFU per 100 g of *Salmonellae*. A treated group of beef steaks was injected with the adduct solution containing HAMMIA. A control group was not injected. Each piece of steak from both groups was left on a tray, sealed with a Saran Wrap™, and stored at 4°C. At four and seven days, the microbial growth in the steaks was measured. At both times there was a 1-4 log reduction in the amount of *E. coli* and *Salmonella* present in the treated group compared to the untreated group.

20 While the preferred compositions or formulations and methods have been disclosed, it will be apparent to those skilled in the art that numerous modifications and variations are possible in light of the above teaching. It should also be realized by those skilled in the art that such modifications and variations do not depart from the spirit and scope of the invention as set forth in
25 the appended claims.